Periareolar (circumareolar) mastopexy with conical breast implants: treatment of ptosis, hypomastia, and changes in position and size of the nipple-areola complex

ABSTRACT

Introduction: Mastopexy surgery associated with breast augmentation is increasingly being requested. At the same time, patients with lower degrees of ptosis are less receptive to any correction through vertical scars. In this context, periareolar (circumareolar) mastopexy with a conical implant is an option that results in a scar that is limited to the areolar perimeter, and enables the treatment of changes in the position and size of the nipple-areola complex. Method: We evaluated 22 patients submitted to periareolar mastopexy with use of conical prostheses coated with polyurethane and placed in a subglandular position. All patients were operated on by the same surgeon. Results: Among the patients, 45% presented with grade I ptosis, 32% grade II, and 23% grade III, and 86% exhibited asymmetry of the nipple-areola complex, 27% lateralization, and 18% large areolas, with some combination in the same patient. The volume of the prostheses ranged from 215 to 380 mL. There were two cases of scar enlargement; however, there were no cases of enlargement of the areola or hypertrophic scar. During the period studied, there were no complications related to the placement of the implant. In the satisfaction questionnaire, most of the patients considered the aspect of the breast to be natural in appearance, giving good grades to the size, symmetry, and height of the breasts; the quality of the scar; and the position, shape, and size of the areolas. Conclusions: This technique produces satisfactory results in cases of mild to moderate ptoses, large areolas, lateralization, and asymmetry of the nipple-areola complex. Important technical aspects include the conservative removal of periareolar skin, use of implants of moderate sizes, and the use of the round-block containment suture.

Keywords: Periareolar; Mastopexy; Ptosis; Implants; Conical; Polyurethane.
INTRODUCTION

Mild to moderate mammary ptosis and changes in the positioning of the nipple-areola complex are frequent findings in patients who wish to undergo breast augmentation surgery. The augmentation mastoplasty associated with mastopexy, especially in these intermediate cases, remains a controversial issue and of great interest to the plastic surgeon because it accounts for the highest rates of complications and postoperative patient dissatisfaction.

The choice of an adequate surgical treatment is crucial because the simple placement of the breast implant without skin removal often results in the recurrence of ptosis. On the other hand, corrections through larger scars, such as vertical incisions, in “L” or inverted “T”, are less accepted by patients who have lower degrees of ptosis and asymmetry of the nipple-areola complex. For these cases, we have used conical breast prostheses together with the removal of periareolar (circumareolar) skin, and round-block dermal sutures. This approach offers a central access to the creation of a secure implant pocket, allows an accurate dissection in all directions, allows repositioning of the nipple-areola complex, and results in a less apparent scar that is camouflaged on the perimeter of the areola.

Several authors have reported on their experiences with periareolar mastopexy and breast augmentation with various types, sizes, and locations of prostheses. However, no studies were found on the use of conical silicone prostheses, which, in theory, could alleviate the problem of breast flattening.

OBJECTIVE

Our aim in this work is to evaluate a series of patients undergoing breast augmentation with periareolar mastopexy by using a polyurethane-coated conical silicone mammary prosthesis placed in a subglandular position. All patients were operated on by the same surgeon.

METHOD

We evaluated 22 patients who underwent breast augmentation with a conical prosthesis, by using periareolar mastopexy with the round-block technique, between February 2011 and February 2013. We studied the following parameters: degree of ptosis, presence of changes in the position and size of the nipple-areola complex, postoperative complications, and patient satisfaction with the surgical outcome.
Surgical Technique (Figures 1 and 2)

With the patient in the standing position, the following markings are placed: middle line (from the sternal notch to the xiphoid process), breast meridians (midclavicular line that passes through the nipple), and inframammary fold. Thereafter, four points of reference are marked. Point A (superior) determines the desired height of the areola, being marked at the intersection of the breast meridian with a height corresponding to the middle third of the arm, normally between 18 and 20 cm from the sternal notch. Points B (distal), C (medial), and D (lateral) are determined by means of digital clamping, with a pinch test. The junction of these points forms a circular or slightly oval figure, which determines the extent of circumareolar skin area to be resected.

With the patient under general anesthesia or epidural anesthesia with sedation, in the 30° supine position, the region to be detached (superiorly at the height of the second intercostal space, medially to about 1 cm from the medial line, laterally on the anterior axillary line, and inferiorly in the inframammary fold) is marked. Marking with a 4-cm-diameter areolatome and the removal of the epidermis marked around the areola are performed. A semicircular infra-areolar incision is made through the breast parenchyma up to the pectoral fascia. The detachment is initially performed digitally (blunt), and then with scissors from the subglandular plane up to the previously defined margins. After the revision of hemostasis, the high or extra-high conical polyurethane-coated prosthesis is placed into the pocket, ensuring that the cone apex remains in place where the nipple-areolar complex is repositioned. The same procedure is repeated in the other breast.

Approximation of the breast tissue and subcutaneous tissue is done with a monocryl 3.0 suture. Then, a circular round-block suture in the deep dermis is done with mononylon 2.0, repositioning the surrounding skin up to the diameter of the 4-cm areolatome. Guidance sutures are placed in the four quadrants with mononylon 4.0. Intradermal suture with monocryl 3.0 is performed in the areolar perimeter. Dressing is applied with gauze, micropore, cotton pad, and compression with crepe bands.

Satisfaction Questionnaire

All patients responded to the satisfaction questionnaire below, with eight questions about the following aspects:

1) Aspect of the breast: natural or artificial.
2) Evaluation with scores of 1 to 4 (1—bad, 2—average, 3—good, or 4—excellent):
   a. Degree of satisfaction with the final result,
   b. Form of the breasts,
   c. Symmetry of the breasts,

Figure 1. (A) Patient marked as described in text. (B) Resection of the circumareolar epidermis. (C) Semicircular infra-areolar incision up to the pectoral fascia. (D) Digital detachment in the subglandular plane. (E) Ample pocket prepared to receive the implant. (F) Placement of the conical prosthesis.

Figure 2. (A) Breasts with implants placed. (B) Approximation of glandular breast tissue with monocryl 3.0. (C) Round-block circular suture with mononylon 2.0 (before approaching the surrounding skin). (D) Right breast already with approximation of the surrounding skin with the round-block suture, up to the diameter of the 4-cm areolatome. (E) Both breasts after the round-block suture. (F) Result after the intradermal suture with monocryl 3.0 in the areolar perimeter.
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d. Height of the breasts,
e. Quality of the scar,
f. Position of the areola,
g. Size of the areola.

RESULTS

The age of the patients at the time of surgery ranged from 22 to 55 years (mean, 33 years). The postoperative follow-up time ranged from 6 to 26 months (mean, 15 months).

Concerning the degree of ptosis, most patients presented with grade I and II ptosis, according to the classification of Regnault. Only five patients had grade III ptosis (Figure 3).

Concerning the presurgery position and the size of the nipple-areola complex, 19 patients exhibited asymmetry, 6 showed lateralization, and 4 had large areolas, with some combination in the same patient (Figure 4).

The volume of prostheses ranged from 215 to 380 mL, with the most frequently used being 300 mL and 345 mL. All prostheses were conical, 5 pairs with a high profile and 18 pairs with an extra-high profile. In four cases, prostheses of different volumes were used to balance differently sized breasts.

Complications related to the placement of the implant were evaluated, such as hematoma, seroma, infection, capsular contracture, undulations, palpable edge, or mobilization of the implant in the period studied. No complications were found.

There were two cases of enlargement of the scar; however, there were no cases of enlargement the areola, hypertrophic scar, or areolar necrosis. In most cases, wrinkling of the periareolar skin was initially observed; however, a complete improvement was observed between 1 and 2 months after the surgery in all cases.

In the satisfaction questionnaire, 21 of 22 patients were satisfied with the results. They considered the aspect of the breast to be natural in appearance (Figure 5), and gave good grades concerning the form, symmetry, and height of the breasts; quality of the scar; and position and size of areolas (Figure 6).

Photographs of the preoperative and postoperative period in some cases are shown in in Figures 7 to 11.
Figure 7. (A, C, E) Preoperative aspect of a 55-year-old patient with grade I ptosis who desired breast augmentation and breast pexy. (B, D, F) Postoperative aspect at 8 months after periareolar mastopexy with a conical prosthesis (300 mL, extra-high profile).

Figure 8. (A, C, E) Preoperative aspect of a 29-year-old patient with grade II ptosis, asymmetry, and sharp lateralization of the nipple-areola complex. (B, D, F) Postoperative aspect at 18 months after periareolar mastopexy with a conical prosthesis (280 mL, high profile).

Figure 9. (A, C, E) Preoperative aspect of a 46-year-old patient with asymmetry and grade I ptosis, who already had a 220-mL breast prosthesis placed 10 years ago and wanted breast augmentation and pexy. (B, D, F) Postoperative aspect at 12 months after periareolar mastopexy and replacement with a conical prosthesis (380 mL, extra-high profile).

Figure 10. (A, C, E) Preoperative aspect of a 33-year-old patient with hypoplasia and grade I mammary ptosis. (B, D, F) Postoperative aspect at 18 months after periareolar mastopexy with a conical prosthesis (380 mL, extra-high profile).
These changes occur with great frequency in patients such as asymmetry, lateralization, or large areolas. Ptosis, other changes of the nipple-areola complex, presented in the preoperative period, in addition to "T", may give better results. The association of a vertical scar, in "L" or inverted "T" indicates that, in cases of more severe ptosis, flattening of the breasts, and poor-quality scars, occurred in up to 40% of cases. In our study, however, we managed to obtain natural-looking breasts, with few complications and a good patient satisfaction index. This significant improvement of results has been reported in other recent studies, and is due to the refinement of the technique and its indications, which will be discussed below.

In the initial work on the use of periareolar mastopexy associated with breast augmentation, a high incidence of unsatisfactory results was reported, mainly owing to the enlargement of the areola, flattening of the breasts, and poor-quality scars, which occurred in up to 40% of cases. In our study, however, we managed to obtain natural-looking breasts, with few complications and a good patient satisfaction index. This significant improvement of results has been reported in other recent studies, and is due to the refinement of the technique and its indications, which will be discussed below.

In relation to the degree of ptosis, most patients (77%) presented with ptosis grades I and II, which, in most of the surgeries, were the main indications for this approach. The three patients who had grade III ptosis had satisfactory results, and this extension of indication has been reported in more recent studies. However, the case with the most advanced grade of ptosis (grade III) was the one that presented scar enlargement and an unsatisfactory result, according to the patient’s own evaluation. This indicates that, in cases of more severe ptosis, the association of a vertical scar, in “L” or inverted “T”, may give better results.

As shown in Figure 4, a large proportion of patients presented in the preoperative period, in addition to ptosis, other changes of the nipple-areola complex, such as asymmetry, lateralization, or large areolas. These changes occur with great frequency in patients and, when small to moderate, can be corrected or softened during the withdrawal of periareolar skin. The correction of these changes in the nipple-areola complex is an important indication of the technique, especially in patients with mild ptosis, in which the simple placement of the implant, in some cases, would correct the ptosis but would not correct asymmetry, lateralization, or large areolas.

The size and shape of the prostheses used plays an important role in the final outcome. As mentioned above, the average volume used was 330 mL (not exceeding 380 mL). This average is in agreement with the literature that shows better results with moderate augmentation; thus, very large volumes should be avoided. More recent studies have used better designed implants, with a high or an extra-high profile, in varying forms, to achieve more natural-looking breasts. In this study, in addition to the use of high- and extra-high-profile prostheses, we suggest the use of conical implants, which has the advantage of presenting a higher projection of the areola and breast, avoiding breast flattening and therefore attaining the most natural result.

In this study, we did not find any complication related to the placement of the implant, such as hematoma, seroma, infection, capsular contracture, undulations, palpable edge, or implant mobilization. We attribute this result to a systematic and careful surgical technique, with blunt juxta muscular dissection just above the fascia, creation of an ample pocket, washing with saline solution, and rigorous hemostasis. In addition, coating the implant with polyurethane may have helped because it was found to be related to a lower incidence of capsular contracture.

Two cases of moderate scar enlargement were observed; however, no cases of hypertrophic scar or change in the form of the areola were observed (the enlargement, flattening, and distortion of the areola are the major complications of the periareolar technique). This result is mainly due to the conservative removal of periareolar skin and the round-block or continuous circumareolar suture technique, in the deep dermis. This suture technique, which controls the tendency of areolar enlargement, allowed for the expansion of the application of various periareolar techniques. In our patients, we used nylon 2.0 suture and a curved needle. Some studies have shown good results with the use of Gore-Tex, a heavy permanent suture that slides well in the dermis, with a straight needle. Other studies propose that the periareolar incision should be made in zigzag form to achieve a more natural-looking final scar. We believe that all improvements are possible; however, the key to controlling the size of the areola is the round-block technique associated with the conservative removal of periareolar skin.

As seen from the questionnaire answered by the patients (Figures 5 and 6), there is a good level of satisfaction with the surgery. From the review of the literature and the experience gained with these surgeries, we conclude that the ideal indications for

**DISCUSSION**

In the initial work on the use of periareolar mastopexy with conical prostheses (extra-high profile, 300 mL in the right breast and 345 mL in the left breast).

**Figure 11.** (A, C, E) Preoperative aspect of a 34-year-old patient with grade III ptosis, mammary asymmetry, and large and lateralized areolas. (B, D, F) Postoperative aspect at 6 months after periareolar mastopexy with conical prostheses (extra-high profile).
breast augmentation with periareolar mastopexy are as follows:

- Large areolas in the preoperative period;
- Mild to moderate nipple and/or breast ptosis that cannot be corrected only with the implant;
- Congenital or acquired asymmetry, for which small adjustments are needed;
- Tuberous breasts;
- Replacement of implants.

Factors favorable to the technique should also be considered, such as breasts with moderate flaccidity, without streaks or skin dystrophies; small to moderate prosthesis volume, up to 380 mL; and elevation of the nipple areola complex <4 cm, preferably up to 2 cm².

We attribute the results obtained in this study to a careful selection of patients, a combination of techniques, and the type of prosthesis used.

**CONCLUSION**

Periareolar (circumareolar) mastopexy with a conical breast prosthesis, by using the round-block suture, allows correcting various common problems, such as mild to moderate ptoses, asymmetries, and changes in the position and size of the nipple areola complex, without major scars or mobilization of breast tissue and resulting in natural-looking breasts with a good degree of patient satisfaction.

**REFERENCES**


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